

09/689,366  
IDS #2



**PCT**

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>6</sup> :</b> <b>C07H 21/04, C12P 21/00, C12Q 1/00</b>	<b>A1</b>	<b>(11) International Publication Number:</b> <b>WO 97/06182</b> <b>(43) International Publication Date:</b> 20 February 1997 (20.02.97)
<b>(21) International Application Number:</b> PCT/US96/12860 <b>(22) International Filing Date:</b> 6 August 1996 (06.08.96)  <b>(30) Priority Data:</b> 08/512,946 8 August 1995 (08.08.95) US 08/569,749 8 December 1995 (08.12.95) US  <b>(71) Applicant:</b> TULARIK, INC. [US/US]; Two Corporate Drive, South San Francisco, CA 94080 (US).  <b>(72) Inventors:</b> ROTHE, Mike; Tularik, Inc., Two Corporate Drive, South San Francisco, CA 94080 (US). GOEDDEL, David, V.; Tularik, Inc., Two Corporate Drive, South San Francisco, CA 94080 (US).  <b>(74) Agents:</b> BREZNER, David, J. et al.; Flehr, Hohbach, Test, Albritton & Herbert, Suite 3400, 4 Embarcadero Center, San Francisco, CA 94111-4187 (US).	<b>(81) Designated States:</b> AU, CA, JP, European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  <b>Published</b> <i>With international search report.</i>	
<b>(54) Title:</b> INHIBITORS OF APOPTOSIS  <b>(57) Abstract</b> <p>The invention provides methods and compositions relating to novel human cellular inhibitor of apoptosis proteins (c-IAP1/2) comprising a series of defined structural domain repeats and/or a RING finger domain; in particular, at least two of: a particular first domain repeat, a particular second domain repeat, and a particular third domain repeat, and/or a particular RING finger domain. The proteins provide a c-IAP specific function, with preferred proteins being capable of modulating the induction of apoptosis; for example, by binding a human tumor necrosis factor receptor associated factor (TRAF). The compositions include nucleic acids which encode the subject c-IAP and hybridization probes and primers capable of hybridizing with the disclosed c-IAP genes. The invention includes methods of using the subject compositions in therapy, in diagnosis and in the biopharmaceutical industry.</p>		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LI	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

*Inhibitors of Apoptosis*

## INTRODUCTION

Field of the Invention

The field of this invention is human proteins involved in the inhibition of apoptosis, or programmed cell death.

5

Background

Cellular apoptosis, or programmed cell death, may be initiated by a variety of different stimuli including viral infection, certain cell-culture conditions, cell-cell signaling, cytokines, etc. Elucidation of signal transduction pathways leading to apoptosis would provide valuable insight into a variety of pathogenic mechanisms. Accordingly, the ability to exogenously modulate the induction of apoptosis would yield therapeutic application for numerous clinical indications. In addition, components of such pathways would provide valuable targets for automated, cost-effective, high throughput drug screening and hence would have immediate application in domestic and international pharmaceutical and biotechnology drug development programs.

15

Relevant Literature

Rothe *et al.* (1994) Cell 78, 681-692, report the existence of tumor necrosis factor (TNF) receptor associated proteins which co-immunoprecipitate with a TNF receptor; see also Rothe, et al., pending US patent application Serial No: 08/446,915. Roy, et al. (1995) Cell 80, 167-178 disclose the gene for a human neuronal apoptosis

20

inhibitory protein. Birnbaum et al. (1994) J Virol 68, 2521-2528 disclose an inhibitor of apoptosis (iap) gene, Op-iap from the Orgyia pseudotsugata nuclear polyhedrosis virus (OpMNPV) with sequence similarity to two other viral genes: Cp-iap derived from Cydia pomonella granulosis virus (CpGV), and iap derived from the Autographa californica nuclear polyhedrosis virus (AcMNPV). Clem and Miller (1994), in Apoptosis II: The Molecular Basis of Apoptosis in Disease, pp 89-110, Cold Spring Harbor Laboratory Press, provide a recent review of apoptosis regulation by insect viruses.

## 10 SUMMARY OF THE INVENTION

The invention provides methods and compositions relating to novel human cellular inhibitor of apoptosis proteins (c-IAP). The subject proteins comprise a series of defined structural domain repeats and/or a RING finger domain; in particular, at least two of a first domain repeat comprising SEQUENCE ID NO: 5 or 6; a second domain repeat comprising SEQUENCE ID NO: 7 or 8; and a third domain repeat comprising SEQUENCE ID NO: 9 or 10; and/or a RING finger domain comprising SEQUENCE ID NO: 11 or 12, or a consensus sequences derived from these human genes. The proteins provide a c-IAP specific function, with preferred proteins being capable of modulating the induction of apoptosis; for example, by binding a human tumor necrosis factor receptor associated factor, TRAF. The compositions include nucleic acids which encode the subject c-IAP and hybridization probes and primers capable of hybridizing with the disclosed c-IAP genes.

The invention includes methods of using the subject compositions in therapy (e.g. gene therapy to enhance expression of a c-IAP gene), in diagnosis (e.g. genetic hybridization screens for c-IAP gene mutations, and in the biopharmaceutical industry (e.g. reagents for increasing yields of recombinant protein by enhancing host cell survival in culture, for screening chemical libraries for lead compounds for a pharmacological agent useful in the diagnosis or treatment of disease associated with apoptosis regulation, etc.).

30

## DETAILED DESCRIPTION OF THE INVENTION

The invention provides methods and compositions relating to novel cellular inhibitor of apoptosis proteins (c-IAPs). The nucleotide sequence of a natural cDNA encoding human c-IAP is shown as SEQUENCE ID NO:1 and the full conceptual translate is shown as SEQUENCE ID NO:2. The nucleotide sequence of another  
5 natural cDNA encoding human c-IAP2 is shown as SEQUENCE ID NO:3 and the full conceptual translate is shown as SEQUENCE ID NO:4. The human c-IAPs of the invention include incomplete translates of SEQUENCE ID NOS:1 and 3 or deletion mutants of SEQUENCE ID NOS: 2 and/or 4, which translates or deletions mutants have at least one of the human c-IAP specific activities described herein. In addition,  
10 the invention provides nonhuman mammalian homologs of the disclosed human c-IAPs. These homologs are encoded by natural cDNAs which are capable of specifically hybridizing with one or more of the disclosed human cDNAs under hybridization conditions describe below and are isolated using the methods and reagents described herein. For example, the amino acid sequence of a murine  
15 homolog of c-IAP1, and the sequence its cDNA are shown in SEQUENCE ID NOS: 14 and 13.

The subject proteins comprise a series of defined structural domain repeats and/or a RING finger domain shown to be necessary for human c-IAP specific function; generally including at least two of: a first domain repeat comprising  
20 SEQUENCE ID NO: 5, 6 or a consensus of 5 and 6, a second domain repeat comprising SEQUENCE ID NO: 7, 8 or a consensus of 7 and 8, and a third domain repeat comprising SEQUENCE ID NO: 9, 10 or a consensus of 9 and 10; and/or a RING finger domain comprising SEQUENCE ID NO: 11, 12 or a consensus of 11 and 12. Preferred domain repeat containing c-IAPs contain each of the three domain  
25 repeats. More preferred c-IAPs comprise the three domain repeats and the C-terminal RING finger. To secure or optimize the requisite function for the protein, the repeats are usually preceded (N-terminally) and separated by intervening regions of about 10 to about 100 residues, which regions preferably derive from those found in the natural c-IAP1 and c-IAP2 translates. Similarly, the RING finger domain of RING finger  
30 domain containing c-IAPs containing proteins is usually preceded by an N-terminal region of about 10 to 300 residues, usually 100 to 300 residues, which region preferably derives from those found in the natural c-IAP1 and c-IAP2 translates.

The proteins provide a human c-IAP1 or c-IAP2 (c-IAP1/2) specific activity or function which may be determined by convenient in vitro, cell-based, or in vivo assays. Preferred proteins are capable of modulating the induction of apoptosis. Such activity or function may be demonstrated in cell culture (e.g. cell transfections) or in animals (e.g. in vivo gene therapy, transgenics). c-IAP1/2 specific function can also be demonstrated by specific binding to a c-IAP1/2 specific binding target, including natural binding targets and nonnatural targets such as c-IAP1/2 specific antibodies. For example, c-IAPs comprising at least two of SEQUENCE ID NOS: 6, 7 and 8 are capable of specifically binding human tumor necrosis factor receptor associated factors 1 and 2 (TRAF1 and TRAF2) in simple in vitro binding assays. Finally, specific function can be assayed immunologically by the ability of the subject protein to elicit a c-IAP1/2 specific antibody in a rodent or rabbit. Generally, human c-IAP1/2-specificity of the binding agent is shown by binding equilibrium constants (usually at least about  $10^7 \text{ M}^{-1}$ , preferably at least about  $10^8 \text{ M}^{-1}$ , more preferably at least about  $10^9 \text{ M}^{-1}$ ). A wide variety of cell-based and cell-free assays may be used to demonstrate human c-IAP1/2-specific binding; preferred are rapid in vitro, cell-free assays such as mediating or inhibiting human c-IAP1/2-protein (e.g. human c-IAP1-TRAF2) binding, immunoassays, etc.

The claimed human c-IAP proteins are isolated, partially pure or pure and are typically recombinantly produced. An "isolated" protein for example, is unaccompanied by at least some of the material with which it is associated in its natural state and constitutes at least about 2%, and preferably at least about 5% by weight of the total protein in a given sample; a partially pure protein constitutes at least about 10%, preferably at least about 30%, and more preferably at least about 60% by weight of the total protein in a given sample; and a pure protein constitutes at least about 70%, preferably at least about 90%, and more preferably at least about 95% by weight of the total protein in a given sample. A wide variety of molecular and biochemical methods are available for generating and expressing the subject compositions, see e.g. Molecular Cloning, A Laboratory Manual (Sambrook, et al. Cold Spring Harbor Laboratory), Current Protocols in Molecular Biology (Eds. Ausubel, et al., Greene Publ. Assoc., Wiley-Interscience, NY) or that are otherwise known in the art.

The invention provides human c-IAP1/2-specific binding agents including substrates, natural intracellular binding targets, etc. and methods of identifying and making such agents, and their use in diagnosis, therapy and pharmaceutical development. For example, human c-IAP1/2-specific agents are useful in a variety of  
5 diagnostic and therapeutic applications, especially where disease or disease prognosis is associated with improper utilization of a pathway involving human c-IAP1/2, e.g. apoptosis. Novel human c-IAP1/2-specific binding agents include human c-IAP1/2-specific antibodies and other natural intracellular binding agents identified with assays such as one- and two-hybrid screens, non-natural intracellular binding agents  
10 identified in screens of chemical libraries, etc.

The invention also provides nucleic acids encoding the subject proteins, which nucleic acids may be part of human c-IAP1/2-expression vectors and may be incorporated into recombinant cells for expression and screening, transgenic animals for functional studies (e.g. the efficacy of candidate drugs for disease associated with  
15 c-IAP1/2 mediated signal transduction), etc., and nucleic acid hybridization probes and replication/amplification primers having a human c-IAP1/2 cDNA specific sequence contained in SEQUENCE ID NO:1 or 3. Nucleic acids encoding human c-IAP1/2 are isolated from eukaryotic cells, preferably human cells, by screening cDNA libraries with probes or PCR primers derived from the disclosed human c-IAP1/2 cDNA.

In addition, the invention provides nucleic acids sharing sufficient sequence similarity with that of the disclosed human c-IAP1/2 cDNAs to effect hybridization thereto. Such human c-IAP1/2 cDNA homologs are capable of hybridizing to the human c-IAP1/2-encoding nucleic acid defined by SEQUENCE ID NO: 1 or 3 under stringency conditions characterized by a hybridization buffer comprising 30%  
25 formamide in 5 x SSPE (0.18 M NaCl, 0.01 M NaPO<sub>4</sub>, pH7.7, 0.001 M EDTA) buffer at a temperature of 42°C and remaining bound when subject to washing at 42°C with the 0.2 x SSPE. Preferred nucleic acids will hybridize in a hybridization buffer comprising 50% formamide in 5 x SSPE buffer at a temperature of 42°C and remain bound when subject to washing at 42°C with 0.2 x SSPE buffer at 42°C. Human  
30 c-IAP1/2 cDNA homologs can also be characterized by BLASTX (Altschul et al. (1990) Basic Local Alignment Search Tool, J Mol Biol 215, 403-410) probability scores. Using this nucleic acid sequence search program BLASTX, complete coding

region human c-IAP1/2 cDNA homologs provide a Probability P(N) score of less than 1.0e-200. More preferred nucleic acids encode c-IAPs with at least about 50%, preferably at least about 60%, more preferably at least 70% pair-wise identity to at least one of SEQUENCE ID NOS: 2 and 4.

- 5           The subject nucleic acids are isolated, i.e. constitute at least about 0.5%, preferably at least about 5% by weight of total nucleic acid present in a given fraction. The subject nucleic acids find a wide variety of applications including use as translatable transcripts, hybridization probes, PCR primers, therapeutic nucleic acids, etc.; use in detecting the presence of human c-IAP1/2 genes and gene transcripts, in  
10   detecting or amplifying nucleic acids encoding additional human c-IAP1/2 homologs and structural analogs, and in gene therapy applications. When used as expression constructs, the nucleic acids are usually recombinant, meaning they comprise a sequence joined to a nucleotide other than that which it is joined to on a natural chromosome. The subject nucleic acids may be contained within vectors, cells or  
15   organisms.

- In diagnosis, c-IAP1/2 hybridization probes find use in identifying wild-type and mutant c-IAP1/2 alleles in clinical and laboratory samples. Mutant alleles are used to generate allele-specific oligonucleotide (ASO) probes for high-throughput clinical diagnoses. In therapy, therapeutic c-IAP1/2 nucleic acids are used to modulate  
20   cellular expression or intracellular concentration or availability of active c-IAP1/2. A wide variety of indications may be treated, either prophylactically or therapeutically with the subject compositions. For example, where cell-specific apoptosis or other limitation of cell growth is desired, e.g. neoproliferative disease, a reduction in c-IAP1/2 expression is effected by introducing into the targeted cell type c-IAP1/2  
25   nucleic acids which reduce the functional expression of c-IAP1/2 gene products (e.g. nucleic acids capable of inhibiting translation of a c-IAP1/2 protein). Conditions for treatment include restenosis, where vascular smooth muscle cells are involved, inflammatory disease states, where endothelial cells, inflammatory cells and glomerular cells are involved, myocardial infarction, where heart muscle cells are  
30   involved, glomerular nephritis, where kidney cells are involved, transplant rejection where endothelial cells are involved, infectious diseases such as HIV infection where certain immune cells and other infected cells are involved, or the like.



These c-IAP1/2 inhibitory nucleic acids are typically antisense: single-stranded sequences comprising complements of the disclosed c-IAP1/2 encoding nucleic acid. Antisense modulation of the expression of a given c-IAP1/2 protein may employ c-IAP1/2 antisense nucleic acids operably linked to gene regulatory sequences. Cell  
5 are transfected with a vector comprising a c-IAP1/2 sequence with a promoter sequence oriented such that transcription of the gene yields an antisense transcript capable of binding to endogenous c-IAP1/2 protein encoding mRNA. Transcription of the antisense nucleic acid may be constitutive or inducible and the vector may provide for stable extrachromosomal maintenance or integration. Alternatively, single-  
10 stranded antisense nucleic acids that bind to genomic DNA or mRNA encoding a given c-IAP1/2 protein may be administered to the target cell, in or temporarily isolated from a host, at a concentration that results in a substantial reduction in expression of the targeted protein.

In other indications, e.g. certain hypersensitivities, atrophic diseases, etc., a  
15 reduction in apoptosis is desired. In these applications, an enhancement in c-IAP1/2 expression is effected by introducing into the targeted cell type c-IAP1/2 nucleic acids which increase the functional expression of c-IAP1/2 gene products. Conditions for treatment include multiple sclerosis, where certain neuronal cells are involved, inflammatory disease states such as rheumatoid arthritis, where bystander cells are  
20 involved, transplant rejection where graft cells are involved, infectious diseases such as HIV infection where certain uninfected host cells are involved, or the like. Such nucleic acids may be c-IAP1/2 expression vectors, vectors which upregulate the functional expression of an endogenous c-IAP1/2 allele, or replacement vectors for targeted correction of c-IAP1/2 mutant alleles.

25 Various techniques may be employed for introducing of the nucleic acids into viable cells. The techniques vary depending upon whether one is using the subject compositions in culture or *in vivo* in a host. Various techniques which have been found efficient include transfection with a retrovirus, viral coat protein-liposome mediated transfection, see Dzaou et al., *Trends in Biotech* 11, 205-210 (1993). In some  
30 situations it is desirable to provide the nucleic acid source with an agent which targets the target cells, such as an antibody specific for a surface membrane protein on the target cell, a ligand for a receptor on the target cell, etc. Where liposomes are

employed, proteins which bind to a surface membrane protein associated with endocytosis may be used for targeting and/or to facilitate uptake, e.g. capsid proteins or fragments thereof tropic for a particular cell type, antibodies for proteins which undergo internalization in cycling, proteins that target intracellular localization and  
5 enhance intracellular half-life. In liposomes, the nucleic acid concentration in the lumen will generally be in the range of about 0.01  $\mu\text{M}$  to 10  $\mu\text{M}$ . For other techniques, the concentration and application rate is determined empirically, using conventional techniques to determine desired ranges.

Application of the subject therapeutics may be systemic or local, i.e.  
10 administered at the site of interest. Various techniques can be used for providing the subject compositions at the site of interest, such as injection, use of catheters, trocars, projectiles, pluronic gel, stents, sustained drug release polymers or other device which provides for internal access. Where an organ or tissue is accessible because of removal from the patient, such organ or tissue may be bathed in a medium containing  
15 the subject compositions, the subject compositions may be painted onto the organ, or may be applied in any convenient way. Systemic administration of the nucleic acid may be effected using naked DNA, lipofection, liposomes with tissue targeting (e.g. antibody).

The invention provides methods and compositions for enhancing the yield of  
20 many recombinantly produced proteins, such as tissue plasminogen activator (t-PA), by increasing maximum cell densities and survival time of host production cells in culture. Specifically, cultured cells are transfected with nucleic acids which effect the up-regulation of endogenous c-IAP or the expression of an exogenous c-IAP. For example, nucleic acids encoding functional c-IAP operably linked to a transcriptional  
25 promoter are used to over-express the exogenous c-IAP in the host cell (see, experimental section, below). Such transformed cells demonstrate enhanced survival ability at elevated cell densities and over extended culture periods.

The invention provides efficient methods of identifying pharmacological agents or lead compounds for agents active at the level of a human c-IAP1/2  
30 modulatable cellular function, particularly human c-IAP1/2 mediated signal transduction, especially in apoptosis. Generally, these screening methods involve assaying for compounds which modulate a human c-IAP1/2 interaction with a natural

c-IAP1/2 binding target. The methods are amenable to automated, cost-effective high throughput screening of chemical libraries for lead compounds. Identified reagents find use in the pharmaceutical industries for animal and human trials; for example, the reagents may be derivatized and rescreened in in vitro and in vivo assays to optimize activity and minimize toxicity for pharmaceutical development. Target indications may include infection, genetic disease, cell growth and regulatory dysfunction, such as neoplasia, inflammation, hypersensitivity, etc.

A wide variety of assays for binding agents are provided including protein-protein binding assays, immunoassays, cell based assays, etc. The human c-IAP1/2 compositions used the methods are usually added in an isolated, partially pure or pure form and are typically recombinantly produced. The human c-IAP1/2 may be part of a fusion product with another peptide or polypeptide, e.g. a polypeptide that is capable of providing or enhancing protein-protein binding, stability under assay conditions (e.g. a tag for detection or anchoring), etc. The assay mixtures comprise a natural intracellular human c-IAP1/2 binding target such as a TRAF. While native binding targets may be used, it is frequently preferred to use portions (e.g. peptides, nucleic acid fragments) thereof so long as the portion provides binding affinity and avidity to the subject human c-IAP1/2 conveniently measurable in the assay. The assay mixture also comprises a candidate pharmacological agent. Candidate agents encompass numerous chemical classes, though typically they are organic compounds; preferably small organic compounds and are obtained from a wide variety of sources including libraries of synthetic or natural compounds. A variety of other reagents may also be included in the mixture. These include reagents like salts, buffers, neutral proteins, e.g. albumin, detergents, etc. which may be used to facilitate optimal binding and/or reduce non-specific or background interactions, etc. Also, reagents that otherwise improve the efficiency of the assay, such as protease inhibitors, nuclease inhibitors, antimicrobial agents, etc. may be used.

The resultant mixture is incubated under conditions whereby, but for the presence of the candidate pharmacological agent, the human c-IAP1/2 specifically binds the cellular binding target, portion or analog with a reference binding affinity. The mixture components can be added in any order that provides for the requisite bindings. Incubations may be performed at any temperature which facilitates optimal

binding, typically between 4 and 40°C, more commonly between 15° and 40°C. Incubation periods are likewise selected for optimal binding but also minimized to facilitate rapid, high-throughput screening, and are typically between .1 and 10 hours, preferably less than 5 hours, more preferably less than 2 hours.

- 5           After incubation, the agent-influenced binding between the human c-IAP1/2 and one or more binding targets is detected by any convenient way. For cell-free binding type assays, a separation step is often used to separate bound from unbound components. Separation may be effected by precipitation (e.g. TCA precipitation, immunoprecipitation, etc.), immobilization (e.g. on a solid substrate), etc., followed by
- 10   washing by, for examples, membrane filtration (e.g. Whatman's P-81 ion exchange paper, Polyfiltronic's hydrophobic GFC membrane, etc.), gel chromatography (e.g. gel filtration, affinity, etc.). In addition, one of the components usually comprises or is coupled to a label. A wide variety of labels may be employed - essentially any label that provides for detection of bound protein. The label may provide for direct
- 15   detection as radioactivity, luminescence, optical or electron density, etc. or indirect detection such as an epitope tag, an enzyme, etc. A variety of methods may be used to detect the label depending on the nature of the label and other assay components. For example, the label may be detected bound to the solid substrate or a portion of the bound complex containing the label may be separated from the solid substrate, and
- 20   thereafter the label detected. Labels may be directly detected through optical or electron density, radiative emissions, nonradiative energy transfers, etc. or indirectly detected with antibody conjugates, etc. For example, in the case of radioactive labels, emissions may be detected directly, e.g. with particle counters or indirectly, e.g. with scintillation cocktails and counters.
- 25           The following experiments and examples are offered by way of illustration and not by way of limitation.

## EXPERIMENTAL

- 30           The murine cellular inhibitor of apoptosis protein 1 (c-IAP1) was biochemically purified as a TNF-R2 associated protein using coimmunoprecipitation Rothe et al. (1994) supra. A large scale protein purification protocol provided material sufficient for peptide sequencing. Fully degenerate oligonucleotides corresponding to

two of the isolated peptides were used to specifically amplify a 0.75 kb DNA fragment from mouse CT6 RNA by Reverse Transcription-PCR. This DNA fragment was used to isolate full-length cDNA clones from a mouse CT6 cDNA library by hybridization (50% formamide, 5xSSPE, 42°C; filters washed at 42°C with 0.2xSSPE, where

5 1xSSPE is 0.18 M NaCl, 0.01 M NaPO<sub>4</sub>, pH7.7, 0.001 M EDTA).

DNA sequence analysis predicted an open reading frame encoding a 612 amino acid protein that shows significant sequence similarity (36 % amino acid identity) with the 'inhibitor of apoptosis protein' (IAP) from insect viruses (Clem, R. J. and Miller, L. K., 1994, supra) and the human 'neuronal apoptosis inhibitory protein' (NAIP) (23 %  
10 amino acid identity), that is involved in spinal muscular atrophy (SMA) an inherited disease in humans (Roy et al., 1995, supra). To obtain the human c-IAP1 gene, the originally amplified mouse DNA fragment was used as a probe to screen a HeLa cDNA library (30% formamide, 5xSSPE, 42°C; filters washed at 42°C with 0.2xSSPE). Sequence analysis of the isolated cDNA clones revealed that they  
15 correspond to two distinct genes, designated c-IAP1 and c-IAP2. The human c-IAP1 cDNA encodes a protein of 618 amino acids that is 84% identical to murine c-IAP1. The human c-IAP2 cDNA encodes a protein of 604 amino acids that shares a high degree of amino acid identity with both murine and human c-IAP1 (72% and 73%, respectively) and represents another member of the IAP superfamily.

20 Comparison of the amino acid sequence of members of the IAP superfamily reveals that they are comprised of at least three distinct domains. The N-terminal region of all IAP family members is comprised of 'baculovirus IAP repeat' (BIR) motifs (Birnbaum et al., 1994, supra). While the viral proteins contain two repeats, the mammalian homologs (c-IAP1, -2) possess three BIR motifs. Similarly, NAIP  
25 contains three BIR repeats. In addition to BIR motifs viral IAPs contain a C-terminal RING finger motif. This Zn-binding domain is also present in c-IAP1 and -2 but not in NAIP. Thus c-IAP1 and -2 define a distinct subfamily within the IAP superfamily that contain three BIR motifs and a RING finger motif. A RING finger domain is also present at the N-terminus of TRAF2 and has been shown to be involved in TRAF2  
30 signal transduction. The RING finger motifs of c-IAP1 and -2 share significant sequence homology with the RING finger domains of viral IAPs but no homology with the TRAF2 RING finger domain besides the conserved cysteine and histidine

residues. The region between the BIR domain and the RING finger domain of c-IAP1 and -2 is strongly conserved but does not reveal any significant homology to other members of the IAP family or any other proteins in the NCBI database.

A yeast two-hybrid system was used to determine how c-IAP1 and -2 interact with TNF-R2 and/or TRAFs. The following results were obtained indistinguishably for c-IAP1 and c-IAP2. Two-hybrid analysis revealed that c-IAP1 does not directly interact with TNF-R2. However, a direct interaction could be detected between c-IAP1 and TRAF2. The conserved TRAF domain of TRAF2 (amino acids 264-501) is sufficient to mediate this interaction. Consistently, c-IAP1 also interacted with TRAF1. Further analysis demonstrated that the coiled-coil region within the TRAF domain of TRAF2 (amino acids 251-358) is required for interaction with c-IAP1. In contrast, the C-terminal region of the TRAF domain (amino acids 359-501) that mediates the association of TNF-R2 with TRAF2 is dispensable for interaction of c-IAP1 with TRAF2. Thus c-IAP1 and TNF-R2 bind to non-overlapping docking sites within the TRAF domain of TRAF2. Consistently, c-IAP1 does not interact with TRAF3 (e.g. Cheng et al. (1995), supra), which does not contain a coiled-coil region with sequence similarity to TRAF2/TRAF1. Deletion mutagenesis of c-IAP1 indicated that the N-terminal half of the protein containing the three BIR motifs (amino acids 1-336 of c-IAP1 and 1-396 of c-IAP2) is sufficient for interaction with TRAF2 and TRAF1. Similarly, combinations of two of the three BIR motifs e.g. amino acid residues 46-99 and 204-249 of c-IAP1 and 29-82 and 189-234 of c-IAP2, separated by IAP1 derived intervening sequences of varying lengths are assayed for TRAF1 and TRAF2 binding. This indicates that BIR motifs represent a novel protein:protein interaction domain. The RING finger domain of c-IAP1/2 (amino acids 571-618 of c-IAP1 and 557-604 of c-IAP2) is not required for interaction with TRAFs, but rather mediates subsequent steps in the c-IAP1/2 signaling cascade. Similarly, a variety of c-IAP1 derived N-terminal leader sequences fused to the c-IAP1 RING finger domain are used to assay signal transduction mediation. In an analogous situation, the RING finger domain of TRAF2 has been demonstrated to be required for TRAF2-mediated activation of NF- $\kappa$ B.

A transfection based co-immunoprecipitation assay was used to investigate how c-IAP1 interacts with the complex of TNF-R2 and TRAFs. In this system c-IAP1

was N-terminally tagged with a FLAG epitope peptide and expressed in human embryonic 293 cells under the control of a constitutive CMV promotor (pRK vector). The c-IAP1 expression vector was transiently co-transfected into 293 cells with expression vectors for TNF-R2 and TRAFs. After 24-36 h, the cells were harvested and extracts immunoprecipitated with anti-TNF-R2 antibodies, followed by Western analysis with anti-FLAG antibodies. This assay demonstrated that while c-IAP1 associates directly with TRAF1 and TRAF2, its interaction with TNF-R2 is indirect and requires the heterocomplex of TRAF1 and TRAF2. Thus, c-IAP1 is a component of the TNF-R2 (CD40)/TRAF signaling complex.

To determine the functional properties of c-IAP1 transient transfection assays were performed in human rhabdomyosarcoma KYM1 cells. The results indicate that overexpression of c-IAP1 but not of control vector, TRAF1 or TRAF2 protects KYM1 cells from TNF-induced programmed cell death (apoptosis). Hence, c-IAP1 regulates the cellular response to TNF by modulating TNF responsiveness, e.g. the initiation of an apoptotic or protective program. The transient transfection assay also finds use as a drug screening assay. In this application, candidate agents are screened as above for their ability to modulate the ability of c-IAP1 to downregulate apoptosis.

#### EXAMPLES

1. Protocol for human c-IAP1 - TRAF2 binding assay.
  - A. Reagents:
    - Neutralite Avidin: 20 µg/ml in PBS.
    - Blocking buffer: 5% BSA, 0.5% Tween 20 in PBS; 1 hour at room temperature.
    - Assay Buffer: 100 mM KCl, 20 mM HEPES pH 7.6, 0.25 mM EDTA, 1% glycerol, 0.5% NP-40, 50 mM β-mercaptoethanol, 1 mg/ml BSA, cocktail of protease inhibitors.
    - <sup>32</sup>P human c-IAP1 10x stock:  $10^{-8}$  -  $10^{-6}$  M unlabeled human c-IAP1 supplemented with 200,000-250,000 cpm of labeled human c-IAP1/21 (Beckman counter). Place in the 4°C microfridge during screening.
    - Protease inhibitor cocktail (1000X): 10 mg Trypsin Inhibitor (BMB # 109894), 10 mg Aprotinin (BMB # 236624), 25 mg Benzamidine (Sigma # B-6506).

25 mg Leupeptin (BMB # 1017128), 10 mg APMSF (BMB # 917575), and 2mM NaVO<sub>3</sub> (Sigma # S-6508) in 10 ml of PBS.

- TRAF2: 10<sup>-4</sup> - 10<sup>-5</sup> M biotinylated truncated TRAF2 (residues 264-501) in PBS.

- 5 B. Preparation of assay plates:
  - Coat with 120 µl of stock N-Avidin per well overnight at 4 °C.
  - Wash 2 times with 200 µl PBS.
  - Block with 150 µl of blocking buffer.
  - Wash 2 times with 200 µl PBS.
- 10 C. Assay:
  - Add 40 µl assay buffer/well.
  - Add 10 µl compound or extract.
  - Add 10 µl <sup>33</sup>P-human c-IAP1 (20,000-25,000 cpm/0.1-10 pmoles/well = 10<sup>-6</sup> - 10<sup>-7</sup> M final concentration).
  - 15
    - Shake at 25°C for 15 minutes.
    - Incubate additional 45 minutes at 25°C.
    - Add 40 µl biotinylated truncated TRAF2 (0.1-10 pmoles/40 ul in assay buffer)
    - Incubate 1 hour at room temperature.
    - 20
      - Stop the reaction by washing 4 times with 200 µl PBS.
      - Add 150 µl scintillation cocktail.
      - Count in Topcount.
  - D. Controls for all assays (located on each plate):
    - a. Non-specific binding
    - 25 b. Soluble (non-biotinylated truncated TRAF2) at 80% inhibition.

All publications and patent applications cited in this specification are herein incorporated by reference as if each individual publication or patent application were specifically and individually indicated to be incorporated by reference. Although the  
30 foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, it will be readily apparent to those of ordinary skill in the art in light of the teachings of this invention that certain changes



and modifications may be made thereto without departing from the spirit or scope of the appended claims.

	SEQUENCE ID NO: 1, 2	h (human) c-IAP1	cDNA, protein
5	SEQUENCE ID NO: 3, 4	h c-IAP2	cDNA, protein
	SEQUENCE ID NO: 5, 6	h c-IAP1,2 repeat 1	protein, protein
	SEQUENCE ID NO: 7, 8	h c-IAP1,2 repeat 2	protein, protein
	SEQUENCE ID NO: 9, 10	h c-IAP1,2 repeat 3	protein, protein
	SEQUENCE ID NO: 11, 12	h c-IAP1,2 RING finger	protein, protein
10	SEQUENCE ID NO: 13, 14	m (murine) c-IAP	cDNA, protein

## SEQUENCE LISTING

## (1) GENERAL INFORMATION:

- (i) APPLICANT: TULARIK, INC.
- (ii) TITLE OF INVENTION: INHIBITORS OF APOPTOSIS
- (iii) NUMBER OF SEQUENCES: 14
- (iv) CORRESPONDENCE ADDRESS:
  - (A) ADDRESSEE: FLEHR, HOHBACH, TEST, ALBRITTON & HERBERT
  - (B) STREET: 4 Embarcadero Center, Suite 3400
  - (C) CITY: San Francisco
  - (D) STATE: California
  - (E) COUNTRY: USA
  - (F) ZIP: 94111
- (v) COMPUTER READABLE FORM:
  - (A) MEDIUM TYPE: Floppy disk
  - (B) COMPUTER: IBM PC compatible
  - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
  - (D) SOFTWARE: PatentIn Release #1.0, Version #1.30
- (vi) CURRENT APPLICATION DATA:
  - (A) APPLICATION NUMBER: PCT/US96/
  - (B) FILING DATE: 06 AUG 1996
  - (C) CLASSIFICATION:
- (vii) PRIOR APPLICATION DATA:
  - (A) APPLICATION NUMBERS: U.S. Serial Nos. 08/512,946 & 08/569,749
  - (B) FILING DATES: 08 AUG 1995 & 08 DEC 1995
  - (C) CLASSIFICATION
- (viii) ATTORNEY/AGENT INFORMATION:
  - (A) NAME: Brezner, David J.
  - (B) REGISTRATION NUMBER: 24,774
  - (C) REFERENCE/DOCKET NUMBER: A-62464/DJB
- (ix) TELECOMMUNICATION INFORMATION:
  - (A) TELEPHONE: (415) 781-1989
  - (B) TELEFAX: (415) 398-3249

## (2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
  - (A) LENGTH: 2589 base pairs
  - (B) TYPE: nucleic acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

TCTAAGTAGT ATCTTGGAAA TTCAGAGAGA TACTCATCCT ACCTGAATAT AAACGAGAT	60
AAATCCAGTA AAGAAAGTGT AGTAAATTCT ACATAAGAGT CTATCATTGA TTTCTTTTGG	120
TGGTAAAAAT CTTAGTTCAT GTGAAGAAAT TTCATGTGAA TGTTTTAGCT ATCAACAGC	180
ACTGTCACCT ACTCATGCAC AAAACTGCCT CCCAAAGACT TTTCCAGGT CCCTCGTATC	240
AAAACATTAA GAGTATAATG GAAGATAGCA CGATCTTGTC AGATTGGACA AACAGCAACA	300

AACAAAAAAT GAAGTATGAC TTTTCCTGTG AACTCTACAG AATGTCTACA TATTCAACTT	360
TCCCCGCCGG GGTGCCTGTC TCAGAAAGGA GTCTTGCTCG TGCTGGTTTT TATTATACTG	420
GTGTGAATGA CAAGGTCAAA TGCTTCTGTT GTGGCCTGAT GCTGGATAAC TGGAAACTAG	480
GAGACAGTCC TATTCAAAAG CATAAACAGC TATATCCTAG CTGTAGCTTT ATTCAGAATC	540
TGSTTTCAGC TAGTCTGGGA TCCACCTCTA AGAATACGTC TCCAATGAGA AACAGTTTTG	600
CACATTCAAT ATCTCCCACC TTGGAACATA GTAGCTTGTT CAGTGGTTCT TACTCCAGCC	660
TTTCTCCAAA CCTCTTAAT TCTAGAGCAG TTGAAGACAT CTCTTCATCG AGGACTAACC	720
CCTACAGTTA TGCAATGAGT ACTGAAGAAG CCAGATTTCT TACCTACCAT ATGTGGCCAT	780
TAACTTTTTT GTCACCATCA GAATTGGCAA GAGCTGGTTT TTATTATATA GGACCTGGAG	840
ATAGGGTAGC CTGCTTTGCC TGTTGGTGGGA AGCTCAGTAA CTGGGAACCA AAGGATGATG	900
CTATGTCAGA ACACCGGAGG CATTTTCCCA ACTGTCCATT TTTGGAAAAT TCTCTAGAAA	960
CTCTGAGGTT TAGCATTTCA AATCTGAGCA TGCAGACACA TGCAGCTCGA ATGAGAACAT	1020
TTATGTACTG GCCATCTAGT GTTCCAGTTC AGCCTGAGCA GCTTGCAAGT GCTGGTTTTT	1080
ATTATGTGGG TCGCAATGAT GATGTCAAAT GCTTTTGTTG TGATGGTGGC TTGAGGTGTT	1140
GGGAATCTGG AGATGATCCA TGGGTAGAAC ATGCCAAGTG GTTTCCAAGG TGTGAGTTCT	1200
TGATACGAAT GAAAGGCCAA GAGTTTGTG ATGAGATTCA AGGTAGATAT CCTCATCTTC	1260
TTGAACAGCT GTTGTCAACT TCAGATACCA CTGGAGAAGA AAATGCTGAC CCACCAATTA	1320
TTCATTTTGG ACCTGGAGAA AGTTCTTCAG AAGATGCTGT CATGATGAAT ACACCTGTGG	1380
TTAAATCTGC CTTGAAATG GGCTTTAATA GAGACCTGGT GAAACAAACA GTTCAAAGTA	1440
AAATCCTGAC AACTGGAGAG AACTATAAAA CAGTTAATGA TATTGTGTCA GCACTTCTAA	1500
ATGCTGAAGA TGAAAAAGA GAGGAGGAGA AGGAAAAACA AGCTGAAGAA ATGGCATCAG	1560
ATGATTTGTC ATTAATTCGG AAGAACAGAA TGGCTCTCTT TCAACAATTG ACATGTGTGC	1620
TTCCTATCCT GGATAATCTT TTAAAGGCCA ATGTAATTAA TAAACAGGAA CATGATATTA	1680
TTAAACAAA AACACAGATA CCTTTACAAG CGAGAGAACT GATTGATACC ATTTTGGTTA	1740
AAGGAAATGC TCGGCCCAAC ATCTTCAAAA ACTGTCTAAA AGAAATTGAC TCTACATTGT	1800
ATAAGAACTT ATTTGTGGAT AAGAAATGA AGTATATTCC AACAGAAGAT GTTTCAGGTC	1860
TGTCACTGGA AGAACAAATG AGGAGGTTGC AAGAAGAACG AACTTGTAAG GTGTGTATGG	1920
ACAAAGAAGT TTCTGTGTGA TTTATTCCTT GTGGTCATCT GGTAGTATGC CAGGAATGTG	1980
CCCCTTCTCT AAGAAAAATG CCTATTGCA GGGGTATAAT CAAGGGTACT GTTCGTACAT	2040
TTCTCTCTTA AAGAAAAATA GTCTATATT TAACCTGCAT AAAAAGGTCT TTAATAATTT	2100
GTTGAACACT TGAAGCCATC TAAAGTAAAA AGGGAATTAT GAGTTTTTCA ATTAGTAACA	2160
TTCATGTTCT AGTCTGCTTT GGTACTAATA ATCTTGTTTC TGAAAAGATG GTATCATATA	2220
TTTAATCTTA ATCTGTTTAT TTACAAGGGA AGATTTATGT TTGGTGAAC ATATTAGTAT	2280
GTATGTGTAC CTAAGGGAGT AGTGTCACTG CTGTTATGC ATCATTTTCTAG GAGTTACTGG	2340

```

ATTTGTTGTT CTTTCAGAAA GCTTTGAATA CTAAATTATA GTGTAGAAAA GAACTGGAAA      2400
CCAGGAAGCTC TGGAGTTCAT CAGAGTTATG GTGCCGAATT GTCTTTGGTG CTTTTCACCTT      2460
GTGTTTAAAA ATAAGGATTI TTCTCTTATT TCTCCCCCTA GTTTGTGAGA AACATCTCAA      2520
TAAAGTGCTT TAAAAA AAAA AAAA AAAA AAAA AAAA AAAA AAAA AAAA AAAA AAAA      2580
AAAAAAA                                           2589

```

## (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 618 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

```

Met His Lys Thr Ala Ser Gln Arg Leu Phe Pro Gly Pro Ser Tyr Gln
 1             5             10             15
Asn Ile Lys Ser Ile Met Glu Asp Ser Thr Ile Leu Ser Asp Trp Thr
      20             25             30
Asn Ser Asn Lys Gln Lys Met Lys Tyr Asp Phe Ser Cys Glu Leu Tyr
      35             40             45
Arg Met Ser Thr Tyr Ser Thr Phe Pro Ala Gly Val Pro Val Ser Glu
      50             55             60
Arg Ser Leu Ala Arg Ala Gly Phe Tyr Tyr Thr Gly Val Asn Asp Lys
      65             70             75             80
Val Lys Cys Phe Cys Cys Gly Leu Met Leu Asp Asn Trp Lys Leu Gly
      85             90             95
Asp Ser Pro Ile Gln Lys His Lys Gln Leu Tyr Pro Ser Cys Ser Phe
      100             105             110
Ile Gln Asn Leu Val Ser Ala Ser Leu Gly Ser Thr Ser Lys Asn Thr
      115             120             125
Ser Pro Met Arg Asn Ser Phe Ala His Ser Leu Ser Pro Thr Leu Glu
      130             135             140
His Ser Ser Leu Phe Ser Gly Ser Tyr Ser Ser Leu Ser Pro Asn Pro
      145             150             155             160
Leu Asn Ser Arg Ala Val Glu Asp Ile Ser Ser Ser Arg Thr Asn Pro
      165             170             175
Tyr Ser Tyr Ala Met Ser Thr Glu Glu Ala Arg Phe Leu Thr Tyr His
      180             185             190
Met Trp Pro Leu Thr Phe Leu Ser Pro Ser Glu Leu Ala Arg Ala Gly
      195             200             205
Phe Tyr Tyr Ile Gly Pro Gly Asp Arg Val Ala Cys Phe Ala Cys Gly
      210             215             220
Gly Lys Leu Ser Asn Trp Glu Pro Lys Asp Asp Ala Met Ser Glu His
      225             230             235             240

```

Arg Arg His Phe Pro Asn Cys Pro Phe Leu Glu Asn Ser Leu Glu Thr  
 245 250 255  
 Leu Arg Phe Ser Ile Ser Asn Leu Ser Met Gln Thr His Ala Ala Arg  
 260 265 270  
 Met Arg Thr Phe Met Tyr Trp Pro Ser Ser Val Pro Val Gln Pro Glu  
 275 280 285  
 Gln Leu Ala Ser Ala Gly Phe Tyr Tyr Val Gly Arg Asn Asp Asp Val  
 290 295 300  
 Lys Cys Phe Cys Cys Asp Gly Gly Leu Arg Cys Trp Glu Ser Gly Asp  
 305 310 315 320  
 Asp Pro Trp Val Glu His Ala Lys Trp Phe Pro Arg Cys Glu Phe Leu  
 325 330 335  
 Ile Arg Met Lys Gly Gln Glu Phe Val Asp Glu Ile Gln Gly Arg Tyr  
 340 345 350  
 Pro His Leu Leu Glu Gln Leu Leu Ser Thr Ser Asp Thr Thr Gly Glu  
 355 360 365  
 Glu Asn Ala Asp Pro Pro Ile Ile His Phe Gly Pro Gly Glu Ser Ser  
 370 375 380  
 Ser Glu Asp Ala Val Met Met Asn Thr Pro Val Val Lys Ser Ala Leu  
 385 390 395 400  
 Glu Met Gly Phe Asn Arg Asp Leu Val Lys Gln Thr Val Gln Ser Lys  
 405 410 415  
 Ile Leu Thr Thr Gly Glu Asn Tyr Lys Thr Val Asn Asp Ile Val Ser  
 420 425 430  
 Ala Leu Leu Asn Ala Glu Asp Glu Lys Arg Glu Glu Glu Lys Glu Lys  
 435 440 445  
 Gln Ala Glu Glu Met Ala Ser Asp Asp Leu Ser Leu Ile Arg Lys Asn  
 450 455 460  
 Arg Met Ala Leu Phe Gln Gln Leu Thr Cys Val Leu Pro Ile Leu Asp  
 465 470 475 480  
 Asn Leu Leu Lys Ala Asn Val Ile Asn Lys Gln Glu His Asp Ile Ile  
 485 490 495  
 Lys Gln Lys Thr Gln Ile Pro Leu Gln Ala Arg Glu Leu Ile Asp Thr  
 500 505 510  
 Ile Leu Val Lys Gly Asn Ala Ala Asn Ile Phe Lys Asn Cys Leu  
 515 520 525  
 Lys Glu Ile Asp Ser Thr Leu Tyr Lys Asn Leu Phe Val Asp Lys Asn  
 530 535 540  
 Met Lys Tyr Ile Pro Thr Glu Asp Val Ser Gly Leu Ser Leu Glu Glu  
 545 550 555 560  
 Gln Leu Arg Arg Leu Gln Glu Glu Arg Thr Cys Lys Val Cys Met Asp  
 565 570 575  
 Lys Glu Val Ser Val Val Phe Ile Pro Cys Gly His Leu Val Val Cys  
 580 585 590

Gln Glu Cys Ala Pro Ser Leu Arg Lys Cys Pro Ile Cys Arg Gly Ile  
           595                                600                                605  
 Ile Lys Gly Thr Val Arg Thr Phe Leu Ser  
           610                                615

## (2) INFORMATION FOR SEQ ID NO:3:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2601 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

```

GGGCAGCAGG TTTACAAAGG AGGAAAACGA CTTCTTCTAG ATTTTITTTT CAGTTTCTTC      60
TATAAATCAA AACTACCTCC CTAGAGAAAG GCTAGTCCCT TTTCTTCCCC ATTCATTTCA      120
TTATGAACAT AGTAGAAAAC AGCATATTCT TATCAAATTT GATGAAAAGC GCCAACACGT      180
TTGAACTGAA ATACGACTTG TCATGTGAAC TGTACCGAAT GTCTACGTAT TCCACTTTTC      240
CTGCTGGGGT CCCTGTCTCA GAAAGGAGTC TTGCTCGCGC TGGTTTCTAT TACACTGGTG      300
TGAATGACAA GGTCAAATGC TTCTGTTGTG GCCTGATGCT GGATAACTGG AAAAGAGGAG      360
ACAGTCCTAC TGAAAAGCAT AAAAAGTTGT ATCCTAGCTG CAGATTCGTT CAGAGTCTAA      420
ATTCCGTTAA CAACTTGGAA GCTACCTCTC AGCCTACTTT TCCTTCTTCA GTAACAAATT      480
CCACACACTC ATTACTTCCG GGTACAGAAA ACAGTGGATA TTTCCGTGGC TCTTATTCAA      540
ACTCTCCATC AAATCCTGTA AACTCCAGAG CAAATCAAGA TTTTCTGCC TTGATGAGAA      600
GTTCTTACCA CTGTGCAATG AATAACGAAA ATGCCAGATT ACTTACTTTT CAGACATGGC      660
CATTGACTTT TCTGTCGCCA ACAGATCTGG CAAAAGCAGG CTTTACTAC ATAGGACCTG      720
GAGACAGAGT GGCTTGCTTT GCCTGTGGTG GAAAATTGAG CAATTGGGAA CCGAAGGATA      780
ATGCTATGTC AGAACACCTG AGACATTTTC CCAAATGCCC ATTTATAGAA AATCAGCTTC      840
AAGACACTTC AAGATACACA GTTTCTAATC TGAGCATGCA GACACATGCA GCCCCTTTA      900
AAACATTCTT TAACTGGCCC TCTAGTGTTT TAGTTAATCC TGAGCAGCTT GCAAGTGCGG      960
GTTTTTATTA TGTGGGTAAC AGTGATGATG TCAAATGCTT TTGCTGTGAT GGTGGACTCA     1020
GGTGTGGGGA ATCTGGAGAT GATCCATGGG TTCAACATGC CAAGTGGTTT CCAAGGTGTG     1080
AGTACTTGAT AAGAATTAAA GGACAGGAGT TCATCCGTCA AGTTCAAGCC AGTTACCCTC     1140
ATCTACTTGA ACAGCTGCTA TCCACATCAG ACAGCCCAGG AGATGAAAAT GCAGAGTCAT     1200
CAATTATCCA TTTTGAACCT GGAGAAGACC ATTCAGAAGA TGCAATCATG ATGAATACTC     1260
CTGTGATTAA TGCTGCCGTG GAAATGGGCT TTAGTAGAAG CCTGGTAAAA CAGACAGTTC     1320
AGAGAAAAAT CCTAGCAACT GGAGAGAATT ATAGACTAGT CAATGATCTT GTGTTAGACT     1380
TACTCAATGC AGAAGATGAA ATAAGGGAAG AGGAGAGAGA AAGAGCAACT GAGGAAAAAG     1440

```

```

AATCAAATGA TTTATTATTA ATCCGGAAGA ATAGAATGGC ACTTTTTCAG CATTTGACTT 1500
GTGTAATTCC AATCCTGGAT AGTCTACTAA CTGCCGGAAT TATTAATGAA CAAGAACATG 1560
ATGTTATTAA ACAGAAGACA CAGACGTCTT TACAAGCAAG AGAACTGATT GATACGATTT 1620
TAGTAAAAGG AAATATTGCA GCCACTGTAT TCAGAAACTC TCTGCAAGAA GCTGAAGCTG 1680
TGTTATATGA GCATTTATTT GTGCAACAGG ACATAAAATA TATTCCCACA GAAGATGTTT 1740
CAGATCTACC AGTGAAGAA CAATTGCGGA GACTACAAGA AGAAAGAACA TGTAAAGTGT 1800
GTATGGACAA AGAAGTGTC ATAGTGTTTA TTCCTTGTTG TCATCTAGTA GTATGCAAG 1860
ATTGTGCTCC TTCTTAAGA AAGTGCTCTA TTTGTAGGAG TACAATCAAG GGTACAGTTC 1920
GTACATTTCT TTCATGAAGA AGAACCAAAA CATCATCTAA ACTTTAGAAT TAATTTATTA 1980
AATGTATTAT AACTTTAACT TTCATCCTAA TTTGGTTTCC TTAATAATTT TATTTATTTA 2040
CAACTCAACA AACATTGTTT TGTGTAACAT ATTTAATATA TGTATCTAAA CCATATGAAC 2100
ATATATTTTT TAGAACTTAA GAGAATGATA GGCTTTTGTT CTTATGAACG AAAAAGAGGT 2160
AGCACTACAA ACACAATATT CAATCAAAAT TTCAGCATT TGAATTTGT AAGTGAAGTA 2220
AAACTTAAGA TATTTGAGTT AACCTTTAAG AATTTTAAAT ATTTTGGCAT TGTACTAATA 2280
CCGGGAACAT GAAGCCAGGT GTGGTGGTAT GTGCCTGTAG TCCCAGGCTG AGGCAAGAGA 2340
ATTACTTGAG CCCAGGAGTT TGAATCCATC CTGGGCAGCA TACTGAGACC CTGCCTTTAA 2400
AAACAAACAG AACAAAAACA AAACACCAGG GACACATTC TCTGTCTTTT TTGATCAGTG 2460
TCCTATACAT CGAAGGTGTG CATATATGTT GAATGACATT TTAGGGACAT GGTGTTTTTA 2520
TAAAGAATTC TGTGAGAAAA AATTTAATAA AACCCCCCAA ATTAAAAAAA AAAAAAAAAA 2580
AAAAAAAAAA AAAAAAAAAA A 2601

```

## (2) INFORMATION FOR SEQ ID NO:4:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 604 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

```

Met Asn Ile Val Glu Asn Ser Ile Phe Leu Ser Asn Leu Met Lys Ser
1           5           10           15
Ala Asn Thr Phe Glu Leu Lys Tyr Asp Leu Ser Cys Glu Leu Tyr Arg
20          25          30
Met Ser Thr Tyr Ser Thr Phe Pro Ala Gly Val Pro Val Ser Glu Arg
35          40          45
Ser Leu Ala Arg Ala Gly Phe Tyr Tyr Thr Gly Val Asn Asp Lys Val
50          55          60
Lys Cys Phe Cys Cys Gly Leu Met Leu Asp Asn Trp Lys Arg Gly Asp
65          70          75          80

```

Ser Pro Thr Glu Lys His Lys Lys Leu Tyr Pro Ser Cys Arg Phe Val  
 85 90 95  
 Gln Ser Leu Asn Ser Val Asn Asn Leu Glu Ala Thr Ser Gln Pro Thr  
 100 105 110  
 Phe Pro Ser Ser Val Thr Asn Ser Thr His Ser Leu Leu Pro Gly Thr  
 115 120 125  
 Glu Asn Ser Gly Tyr Phe Arg Gly Ser Tyr Ser Asn Ser Pro Ser Asn  
 130 135 140  
 Pro Val Asn Ser Arg Ala Asn Gln Asp Phe Ser Ala Leu Met Arg Ser  
 145 150 155 160  
 Ser Tyr His Cys Ala Met Asn Asn Glu Asn Ala Arg Leu Leu Thr Phe  
 165 170 175  
 Gln Thr Trp Pro Leu Thr Phe Leu Ser Pro Thr Asp Leu Ala Lys Ala  
 180 185 190  
 Gly Phe Tyr Tyr Ile Gly Pro Gly Asp Arg Val Ala Cys Phe Ala Cys  
 195 200 205  
 Gly Gly Lys Leu Ser Asn Trp Glu Pro Lys Asp Asn Ala Met Ser Glu  
 210 215 220  
 His Leu Arg His Phe Pro Lys Cys Pro Phe Ile Glu Asn Gln Leu Gln  
 225 230 235 240  
 Asp Thr Ser Arg Tyr Thr Val Ser Asn Leu Ser Met Gln Thr His Ala  
 245 250 255  
 Ala Arg Phe Lys Thr Phe Phe Asn Trp Pro Ser Ser Val Leu Val Asn  
 260 265 270  
 Pro Glu Gln Leu Ala Ser Ala Gly Phe Tyr Tyr Val Gly Asn Ser Asp  
 275 280 285  
 Asp Val Lys Cys Phe Cys Cys Asp Gly Gly Leu Arg Cys Trp Glu Ser  
 290 295 300  
 Gly Asp Asp Pro Trp Val Gln His Ala Lys Trp Phe Pro Arg Cys Glu  
 305 310 315 320  
 Tyr Leu Ile Arg Ile Lys Gly Gln Glu Phe Ile Arg Gln Val Gln Ala  
 325 330 335  
 Ser Tyr Pro His Leu Leu Glu Gln Leu Leu Ser Thr Ser Asp Ser Pro  
 340 345 350  
 Gly Asp Glu Asn Ala Glu Ser Ser Ile Ile His Phe Glu Pro Gly Glu  
 355 360 365  
 Asp His Ser Glu Asp Ala Ile Met Met Asn Thr Pro Val Ile Asn Ala  
 370 375 380  
 Ala Val Glu Met Gly Phe Ser Arg Ser Leu Val Lys Gln Thr Val Gln  
 385 390 395 400  
 Arg Lys Ile Leu Ala Thr Gly Glu Asn Tyr Arg Leu Val Asn Asp Leu  
 405 410 415  
 Val Leu Asp Leu Leu Asn Ala Glu Asp Glu Ile Arg Glu Glu Glu Arg  
 420 425 430



Glu Arg Ala Thr Glu Glu Lys Glu Ser Asn Asp Leu Leu Leu Ile Arg  
 435 440 445  
 Lys Asn Arg Met Ala Leu Phe Gln His Leu Thr Cys Val Ile Pro Ile  
 450 455 460  
 Leu Asp Ser Leu Leu Thr Ala Gly Ile Ile Asn Glu Gln Glu His Asp  
 465 470 475 480  
 Val Ile Lys Gln Lys Thr Gln Thr Ser Leu Gln Ala Arg Glu Leu Ile  
 485 490 495  
 Asp Thr Ile Leu Val Lys Gly Asn Ile Ala Ala Thr Val Phe Arg Asn  
 500 505 510  
 Ser Leu Gln Glu Ala Glu Ala Val Leu Tyr Glu His Leu Phe Val Gln  
 515 520 525  
 Gln Asp Ile Lys Tyr Ile Pro Thr Glu Asp Val Ser Asp Leu Pro Val  
 530 535 540  
 Glu Glu Gln Leu Arg Arg Leu Gln Glu Glu Arg Thr Cys Lys Val Cys  
 545 550 555 560  
 Met Asp Lys Glu Val Ser Ile Val Phe Ile Pro Cys Gly His Leu Val  
 565 570 575  
 Val Cys Lys Asp Cys Ala Pro Ser Leu Arg Lys Cys Pro Ile Cys Arg  
 580 585 590  
 Ser Thr Ile Lys Gly Thr Val Arg Thr Phe Leu Ser  
 595 600

## (2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 55 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

Cys Glu Leu Tyr Arg Met Ser Thr Tyr Ser Thr Phe Pro Ala Gly Val  
 1 5 10 15  
 Pro Val Ser Glu Arg Ser Leu Ala Arg Ala Gly Phe Tyr Tyr Thr Gly  
 20 25 30  
 Val Asn Asp Lys Val Lys Cys Phe Cys Cys Gly Leu Met Leu Asp Asn  
 35 40 45  
 Trp Lys Leu Gly Asp Ser Pro  
 50 55

## (2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 55 amino acids
  - (B) TYPE: amino acid
  - (C) STRANDEDNESS: single
  - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

```

Cys Glu Leu Tyr Arg Met Ser Thr Tyr Ser Thr Phe Pro Ala Gly Val
1           5           10           15
Pro Val Ser Glu Arg Ser Leu Ala Arg Ala Gly Phe Tyr Tyr Thr Gly
          20           25           30
Val Asn Asp Lys Val Lys Cys Phe Cys Cys Gly Leu Met Leu Asp Asn
          35           40           45
Trp Lys Arg Gly Asp Ser Pro
          50           55

```

(2) INFORMATION FOR SEQ ID NO:7:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 46 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

```

Leu Ala Arg Ala Gly Phe Tyr Tyr Ile Gly Pro Gly Asp Arg Val Ala
1           5           10           15
Cys Phe Ala Cys Gly Gly Lys Leu Ser Asn Trp Glu Pro Lys Asp Asp
          20           25           30
Ala Met Ser Glu His Arg Arg His Phe Pro Asn Cys Pro Phe
          35           40           45

```

(2) INFORMATION FOR SEQ ID NO:8:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 46 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

```

Leu Ala Lys Ala Gly Phe Tyr Tyr Ile Gly Pro Gly Asp Arg Val Ala
1           5           10           15
Cys Phe Ala Cys Gly Gly Lys Leu Ser Asn Trp Glu Pro Lys Asp Asn
          20           25           30
Ala Met Ser Glu His Leu Arg His Phe Pro Lys Cys Pro Phe
          35           40           45

```

(2) INFORMATION FOR SEQ ID NO:9:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 48 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single

(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

```

Pro Glu Gln Leu Ala Ser Ala Gly Phe Tyr Tyr Val Gly Arg Asn Asp
1           5           10           15
Asp Val Lys Cys Phe Cys Cys Asp Gly Gly Leu Arg Cys Trp Glu Ser
          20           25           30
Gly Asp Asp Pro Trp Val Glu His Ala Lys Trp Phe Pro Arg Cys Glu
          35           40           45

```

(2) INFORMATION FOR SEQ ID NO:10:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 48 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

```

Pro Glu Gln Leu Ala Ser Ala Gly Phe Tyr Tyr Val Gly Asn Ser Asp
1           5           10           15
Asp Val Lys Cys Phe Cys Cys Asp Gly Gly Leu Arg Cys Trp Glu Ser
          20           25           30
Gly Asp Asp Pro Trp Val Gln His Ala Lys Trp Phe Pro Arg Cys Glu
          35           40           45

```

(2) INFORMATION FOR SEQ ID NO:11:

(i) SEQUENCE CHARACTERISTICS:  
 (A) LENGTH: 52 amino acids  
 (B) TYPE: amino acid  
 (C) STRANDEDNESS: single  
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

```

Glu Glu Arg Thr Cys Lys Val Cys Met Asp Lys Glu Val Ser Val Val
1           5           10           15
Phe Ile Pro Cys Gly His Leu Val Val Cys Gln Glu Cys Ala Pro Ser
          20           25           30
Leu Arg Lys Cys Pro Ile Cys Arg Gly Ile Ile Lys Gly Thr Val Arg
          35           40           45
Thr Phe Leu Ser
          50

```

## (2) INFORMATION FOR SEQ ID NO:12:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 52 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

```

Glu Glu Arg Thr Cys Lys Val Cys Met Asp Lys Glu Val Ser Ile Val
1          5          10          15
Phe Ile Pro Cys Gly His Leu Val Val Cys Lys Asp Cys Ala Pro Ser
20          25          30
Leu Arg Lys Cys Pro Ile Cys Arg Ser Thr Ile Lys Gly Thr Val Arg
35          40          45
Thr Phe Leu Ser
50

```

## (2) INFORMATION FOR SEQ ID NO:13:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2862 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: cDNA

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

```

TTCCTTTACA GTGAATACTG TAGTCTTAAT AGACCTGAGC TGACTGCTGC AGTTGATGTA      60
ACCCACTTTA GAGAATACTG TATGACATCT TCTCTAAGGA AAACCAGCTG CAGACTTCAC      120
TCAGTTCCTT TCATTTCATA GGAAAAGGAG TAGTTCAGAT GTCATGTTTA AGTCCTTATA      180
AGGGAAAAGA GCCTGAATAT ATGCCCTAGT ACCTAGGCTT CATAACTAGT AATAAGAAGT      240
TAGTTATGGG TAAATAGATC TCAGGTTACC CAGAAGAGTT CATGTGACCC CCAAAGAGTC      300
CTAACTAGTG TCTTGCCAAG TGAGACAGAT TTGTCCTGTG AGGGTGTCAA TTCACCAAGTC      360
CAAGCAGAAG ACAATGAATC TATCCAGTCA GGTGTCGTGT GTGGAGATCT AGTGTCAAGT      420
GGTGAGAAAC TTCATCTGGA AGTTTAAGCG GTCAGAAATA CTATTACTAC TCATGGACAA      480
AACTGTCTCC CAGAGACTCG GCCAAGGTAC CTTACACCAA AAAGTTAAAC GTATAATGGA      540
GAAGAGCACA ATCTTGTCAG ATTGGACAAA GGAGAGCGAA GAAAAAATGA AGTTTGACTT      600
TTCGTGTGAA CTCTACCGAA TGTCTACATA TTCAGCTTTT CCCAGGGGAG TTCCTGTCTC      660
AGAGAGGAGT CTGGCTCGTG CTGGCTTTTA TTATACAGGT GTGAATGACA AAGTCAAGTG      720
CTTCTGCTGT GGCCTGATGT TGGATAACTG GAAACAAGGG GACAGTCCTG TTGAAAAGCA      780
CAGACAGTTC TATCCCAGCT GCAGCTTTGT ACAGACTCTG CTTTCAGCCA GTCTGCAGTC      840
TCCATCTAAG AATATGTCTC CTGTGAAAAG TAGATTTGCA CATTCGTCAC CTCTGGAACG      900

```

AGGTGGCATT	CACTCCAACC	TGTGCTCTAG	CCCTCTTAAT	TCTAGAGCAG	TGGAAGACTT	960
CTCATCAAGG	ATGGATCCCT	GCAGCTATGC	CATGAGTACA	GAAGAGGCCA	GATTTCTTAC	1020
TTACAGTATG	TGGCCTTTAA	GTTTCTGTG	ACCAGCAGAG	CTGCCCAGAG	CTGGCTTCTA	1080
TTACATAGGG	CCTGGAGACA	GGGTGGCCTG	TTTTGCCTGT	GGTGGGAAAC	TGAGCAACTG	1140
GGAACCAAAG	GATGATGCTA	TGTCAGAGCA	CCGCAGACAT	TTTCCCCACT	GTCCATTTCT	1200
GGAAAATACT	TCAGAAACAC	AGAGGTTTAG	TATATCAAAT	CTAAGTATGC	AGACACACTC	1260
TGCTCGATTG	AGGACATTTT	TGTACTGGCC	ACCTAGTGTT	CCTGTTGAGC	CCGAGCAGCT	1320
TGCAAGTGCT	GGATTCTATT	ACGTGGATCG	CAATGATGAT	GTCAAGTGCT	TTTGTTGTGA	1380
TGGTGGCTTG	AGATGTTGGG	AACCTGGAGA	TGACCCCTGG	ATAGAACACG	CCAAATGGTT	1440
TCCAAGGTGT	GAGTTCTTGA	TACGGATGAA	GGGTCAGGAG	TTTGTTGATG	AGATTCAAGC	1500
TAGATATCCT	CATCTTCTTG	AGCAGCTGTT	GTCCACTTCA	GACACCCAG	GAGAAGAAAA	1560
TGCTGACCCT	ACAGAGACAG	TGGTGCAATT	TGGCCCTGGA	GAAAGTTCGG	AAGATGTCGT	1620
CATGATGAGC	ACGCCCTGTG	TTAAAGCAGC	CTTGGAATG	GGCTTCAGTA	GGAGCCTGGT	1680
GAGACAGACG	GTTACGCGGC	AGATCCTGGC	CACTGGTGAG	AACTACAGGA	CCGTCAATGA	1740
TATTGTCTCA	GTACTTTTGA	ATGCTGAAGA	TGAGAGAAGA	GAAGAGGAGA	AGGAAGACA	1800
GACTGAAGAG	ATGGCATCAG	GTGACTTATC	ACTGATTCGG	AAGAATAGAA	TGGCCCTCTT	1860
TCAACAGTTG	ACACATGTCC	TTCTATCCT	GGATAATCTT	CITGAGGCCA	GTGTAATTAC	1920
AAAACAGGAA	CATGATATTA	TTAGACAGAA	AACACAGATA	CCCTTACAAG	CAAGAGAGCT	1980
TATTGACACC	GTTTTAGTCA	AGGGAAATGC	TGCAGCCAAC	ATCTTCAAAA	ACTCTCTGAA	2040
GGAAATTGAC	TCCACGTTAT	ATGAAAACCT	ATTTGTGGAA	AAGAATATGA	AGTATATTCC	2100
AACAGAAGAC	GTTTCAGGCT	TGTCATTGGA	AGAGCAGTTG	CGGAGATTAC	AAGAAGAACG	2160
AACTTGCAAA	GTGTGTATGG	ACAGAGAGGT	TTCTATTGTG	TTCATTCCGT	GTGGTCATCT	2220
AGTAGTCTGC	CAGGAATGTG	CCCCTTCTCT	AAGGAAGTGC	CCCATCTGCA	GGGGGACAAT	2280
CAAGGGGACT	GTGCGCACAT	TTCTCTCATG	AGTGAAGAAT	GGTCTGAAAG	TATTGTTGGA	2340
CATCAGAAGC	TGTCAGAACA	AAGAATGAAC	TACTGATTTT	AGCTCTTCAG	CAGGACATTC	2400
TACTCTCTTT	CAAGATTAGT	AATCTTGCTT	TATGAAGGGT	AGCATTGTAT	ATTTAAGCTT	2460
AGTCTGTGTC	AAGGGAAGGT	CTATGCTGTT	GAGCTACAGG	ACTGTGTCTG	TTCCAGAGCA	2520
GGAGTTGGGA	TGCTTGCTGT	ATGTCCTTCA	GGACTTCTTG	GATTTGGAAT	TTGTGAAAGC	2580
TTTGGAATTCA	GGTGATGTGG	AGCTCAGAAA	TCCTGAAACC	AGTGGCTCTG	GTAATCAGTA	2640
GTTAGGGTAC	CCTGTGCTTC	TTGGTGCTTT	TCCTTTCTGG	AAAATAAGGA	TTTTTCTGCT	2700
ACTGGTAAAT	ATTTTCTGTT	TGTGAGAAAT	ATATTAAAGT	GTTTCTTTTA	AAGGCGTGCA	2760
TCATTGTAGT	GTGTGCAGGG	ATGTATGCAG	GCAAAACACT	GTGTATATAA	TAAATAAATC	2820
TTTTTAAAAA	GTGAAAAAAA	AAAAAAAAAA	AAAAAAAAAA	AA		2862

## (2) INFORMATION FOR SEQ ID NO:14:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 612 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

```

Met Asp Lys Thr Val Ser Gln Arg Leu Gly Gln Gly Thr Leu His Gln
 1           5          10          15
Lys Leu Lys Arg Ile Met Glu Lys Ser Thr Ile Leu Ser Asn Trp Thr
 20          25          30
Lys Glu Ser Glu Glu Lys Met Lys Phe Asp Phe Ser Cys Glu Leu Tyr
 35          40          45
Arg Met Ser Thr Tyr Ser Ala Phe Pro Arg Gly Val Pro Val Ser Glu
 50          55          60
Arg Ser Leu Ala Arg Ala Gly Phe Tyr Tyr Thr Gly Val Asn Asp Lys
 65          70          75          80
Val Lys Cys Phe Cys Cys Gly Leu Met Leu Asp Asn Trp Lys Gln Gly
 85          90          95
Asp Ser Pro Val Glu Lys His Arg Gln Phe Tyr Pro Ser Cys Ser Phe
100          105          110
Val Gln Thr Leu Leu Ser Ala Ser Leu Gln Ser Pro Ser Lys Asn Met
115          120          125
Ser Pro Val Lys Ser Arg Phe Ala His Ser Ser Pro Leu Glu Arg Gly
130          135          140
Gly Ile His Ser Asn Leu Cys Ser Ser Pro Leu Asn Ser Arg Ala Val
145          150          155          160
Glu Asp Phe Ser Ser Arg Met Asp Pro Cys Ser Tyr Ala Met Ser Thr
165          170          175
Glu Glu Ala Arg Phe Leu Thr Tyr Ser Met Trp Pro Leu Ser Phe Leu
180          185          190
Ser Pro Ala Glu Leu Ala Arg Ala Gly Phe Tyr Tyr Ile Gly Pro Gly
195          200          205
Asp Arg Val Ala Cys Phe Ala Cys Gly Gly Lys Leu Ser Asn Trp Glu
210          215          220
Pro Lys Asp Asp Ala Met Ser Glu His Arg Arg His Phe Pro His Cys
225          230          235          240
Pro Phe Leu Glu Asn Thr Ser Glu Thr Gln Arg Phe Ser Ile Ser Asn
245          250          255
Leu Ser Met Gln Thr His Ser Ala Arg Leu Arg Thr Phe Leu Tyr Trp
260          265          270
Pro Pro Ser Val Pro Val Gln Pro Glu Gln Leu Ala Ser Ala Gly Phe
275          280          285

```

Tyr Tyr Val Asp Arg Asn Asp Asp Val Lys Cys Phe Cys Cys Asp Gly  
 290 295 300  
 Gly Leu Arg Cys Trp Glu Pro Gly Asp Asp Pro Trp Ile Glu His Ala  
 305 310 315 320  
 Lys Trp Phe Pro Arg Cys Glu Phe Leu Ile Arg Met Lys Gly Gln Glu  
 325 330 335  
 Phe Val Asp Glu Ile Gln Ala Arg Tyr Pro His Leu Leu Glu Gln Leu  
 340 345 350  
 Leu Ser Thr Ser Asp Thr Pro Gly Glu Glu Asn Ala Asp Pro Thr Glu  
 355 360 365  
 Thr Val Val His Phe Gly Pro Gly Glu Ser Ser Glu Asp Val Val Met  
 370 375 380  
 Met Ser Thr Pro Val Val Lys Ala Ala Leu Glu Met Gly Phe Ser Arg  
 385 390 395 400  
 Ser Leu Val Arg Gln Thr Val Gln Arg Gln Ile Leu Ala Thr Gly Glu  
 405 410 415  
 Asn Tyr Arg Thr Val Asn Asp Ile Val Ser Val Leu Leu Asn Ala Glu  
 420 425 430  
 Asp Glu Arg Arg Glu Glu Glu Lys Glu Arg Gln Thr Glu Glu Met Ala  
 435 440 445  
 Ser Gly Asp Leu Ser Leu Ile Arg Lys Asn Arg Met Ala Leu Phe Gln  
 450 455 460  
 Gln Leu Thr His Val Leu Pro Ile Leu Asp Asn Leu Leu Glu Ala Ser  
 465 470 475 480  
 Val Ile Thr Lys Gln Glu His Asp Ile Ile Arg Gln Lys Thr Gln Ile  
 485 490 495  
 Pro Leu Gln Ala Arg Glu Leu Ile Asp Thr Val Leu Val Lys Gly Asn  
 500 505 510  
 Ala Ala Ala Asn Ile Phe Lys Asn Ser Leu Lys Glu Ile Asp Ser Thr  
 515 520 525  
 Leu Tyr Glu Asn Leu Phe Val Glu Lys Asn Met Lys Tyr Ile Pro Thr  
 530 535 540  
 Glu Asp Val Ser Gly Leu Ser Leu Glu Glu Gln Leu Arg Arg Leu Gln  
 545 550 555 560  
 Glu Glu Arg Thr Cys Lys Val Cys Met Asp Arg Glu Val Ser Ile Val  
 565 570 575  
 Phe Ile Pro Cys Gly His Leu Val Val Cys Gln Glu Cys Ala Pro Ser  
 580 585 590  
 Leu Arg Lys Cys Pro Ile Cys Arg Gly Thr Ile Lys Gly Thr Val Arg  
 595 600 605  
 Thr Phe Leu Ser  
 610

WHAT IS CLAIMED IS:

1. An isolated nucleic acid having the sequence defined by SEQUENCE ID NO: 1 or 3, or a fragment thereof capable of specifically hybridizing with a nucleic acid  
5 having the sequence defined by SEQUENCE ID NO: 1 or 3 under stringency conditions defined by a hybridization buffer comprising 30% formamide in 5 x SSPE (0.18 M NaCl, 0.01 M NaPO<sub>4</sub>, pH7.7, 0.001 M EDTA) buffer at a temperature of 42°C and remaining bound when subject to washing at 42°C with the 0.2 x SSPE.
- 10 2. An isolated nucleic acid according to claim 1 capable of specifically hybridizing with a nucleic acid having the sequence defined by SEQUENCE ID NO: 1 or 3 under stringency conditions defined by a hybridization buffer comprising 50% formamide in 5 x SSPE buffer at a temperature of 42°C and remaining bound when subject to washing at 42°C with 0.2 x SSPE buffer at 42°C.
- 15 3. An isolated nucleic acid according to claim 1 encoding a human cellular inhibitor of apoptosis protein (c-IAP) comprising at least two of: a first domain comprising SEQUENCE ID NO: 5 or 6, a second domain comprising SEQUENCE ID NO: 7 or 8, and a third domain comprising SEQUENCE ID NO: 9 or 10; said protein  
20 having a c-IAP specific activity.
4. A method of making a human cellular inhibitor of apoptosis protein (c-IAP) comprising introducing a nucleic acid according to claim 3 into a host cell, growing said host cell under conditions whereby said nucleic acid is expressed as a transcript  
25 and said transcript is expressed as a translation product comprising a cellular inhibitor of apoptosis protein, and isolating said translation product.
5. A method of identifying lead compounds for a pharmacological agent useful in the diagnosis or treatment of disease, said method comprising the steps of:  
30 incubating a mixture comprising:  
a human c-IAP made by a method according to claim 4,



a natural intracellular human c-IAP binding target, wherein said binding target is capable of specifically binding said human c-IAP, and a candidate pharmacological agent;

under conditions whereby, but for the presence of said candidate

5 pharmacological agent, said human c-IAP specifically binds said binding target at a reference affinity;

detecting the binding affinity of said human c-IAP to said binding target to determine an agent-biased affinity,

wherein a difference between the agent-biased affinity and the test affinity

10 indicates that said candidate pharmacological agent is a lead compound for a pharmacological agent capable of modulating human c-IAP-dependent signal transduction.

6. A method according to claim 5, wherein said human c-IAP binding target  
15 comprises a TRAF or an intracellular fragment of a TRAF sufficient to provide for c-IAP-specific binding.

7. A method of modulating apoptosis regulation in a cell comprising introducing  
into said cell a nucleic acid according to claim 1 whereby said nucleic acid is  
20 expressed in said cell and the resultant gene product modulates apoptosis regulation in said cell.

8. A method of modulating apoptosis regulation in a cell comprising introducing  
into said cell a nucleic acid according to claim 3 whereby said nucleic acid is  
25 expressed in said cell and the resultant gene product modulates apoptosis regulation in said cell.

9. A method according to claim 8 wherein said cell expresses a recombinant  
protein in in vitro culture and said gene product inhibits apoptosis in said cell,  
30 whereby the yield of said recombinant protein is increased.

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US96/12860**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(6) : CO7H 21/04; C12P 21/00; C12Q 1/00

US CL : 435/5, 69.1; 536/23.1

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/5, 69.1; 536/23.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

STN: Medline, Biosis, CAPUS, WPIDS  
apoptosis, iap(s), TRAF(s)**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X,P	ROTHE et al. The TNFR2-TRAF signaling complex contains two novel proteins related to baculoviral inhibitor of apoptosis proteins. Cell. 29 December 1995, Vol.83, pages 1243-1252, see entire document.	1-9
A,P	CLEM et al. Anti-apoptotic genes of baculoviruses. Cell death and differentiation. January 1996, Vol.3, pages 9-16.	1-9
A	ROY et al. The gene for neuronal apoptosis inhibitory protein is partially deleted in individuals with spinal muscular atrophy. Cell. 13 January 1995, Vol.80, pages 167-178.	1-9

☒ Further documents are listed in the continuation of Box C.
 ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"G" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

11 SEPTEMBER 1996

Date of mailing of the international search report

24 OCT 1996

Name and mailing address of the ISA/US  
Commissioner of Patents and Trade Marks  
Box PCT  
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

ROBERT A. SCHWARTZMAN

Telephone No. (703) 308-0196

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US96/12860

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	ROTHE et al. TRAF2-mediated activation of NF-kB by TNF receptor 2 and CD40. Science.08 September 1995, Vol.269, pages 1424-1427.	1-9